

SQS as a conformity assessment body identification number 1250 herewith certifies the company

**INMATEC Gase
Technologie GmbH & Co. KG
Gewerbstrasse 72
82211 Herrsching
Germany**

the use of a quality assurance system in its design, development, manufacturing and distribution which fulfills the requirements set out in:

ANNEX II

Directive 93/42/EEC (without section 4)

This approval is based on the report dated March 20, 2019.

The scope of validity covers the products

**Medical oxygensgenerators of the type
IMT-POC 8000 to 8950 med**

The following CE label can be applied to these products mentioned in the Appendix of this certificate

CE 1250

A condition for the validity of this certificate is a regular examination in accordance with Annex II.5 of the Directive 93/42/EEC.

Validity 22.03.2019–21.03.2022
Issue 22.03.2019

Reg. no. 36160
Approved Medical Responsible
22.03.2019



F. Müller, CEO SQS



D. Taddeo, Medical Responsible



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ANNEX II

Directive 93/42/EEC (without section 4)

This Appendix is valid only in connection with the following certificate:

Registration Number 36160

Validity from March 22, 2019 up to and including March 21, 2022

This approval includes the following Medical Device/s:

Class IIa

Oxygensgenerators of type:

IMT-POC 8150 med

IMT-POC 8250 med

IMT-POC 8350 med

IMT-POC 8450 med

IMT-POC 8550 med

IMT-POC 8650 med

IMT-POC 8000 med

IMT-POC 8100 med

IMT-POC 8200 med

IMT-POC 8300 med

IMT-POC 8400 med

IMT-POC 8500 med

IMT-POC 8600 med

IMT-POC 8700 med

IMT-POC 8800 med

IMT-POC 8900 med

IMT-POC 8910 med

IMT-POC 8920 med

IMT-POC 8930 med

IMT-POC 8940 med

IMT-POC 8950 med

Appendix issued: 22.03.2019

